

Minutes (Draft)  
Scientific Advisory Committee Meeting  
January 8, 2008  
DFS Central Laboratory, Classroom 1

Committee Members Present:

Dr. Frederick Bieber  
Mr. Joseph Bono  
Dr. Dale Carpenter  
Mr. Dominic Denio  
Dr. Arthur Eisenberg  
Mr. Barry Fisher, Chair  
Ms. Deborah Friedman  
Dr. Dan Krane (via teleconference)  
Mr. Pete Marone  
Dr. Alphonse Poklis  
Dr. Norah Rudin  
Mr. Kenneth Smith

Committee Member Absent:

Dr. Jose Almirall

Staff Members Present

Ms. Wanda Adkins, Office Manager  
Mr. Jeffrey Ban, Central Laboratory Director  
Dr. David Barron, Technical Services Director  
Ms. Eileen Davis, Trace Section Chief  
Ms. Leslie Ellis, Human Resource Manager  
Ms. Michele Gowdy, Department Counsel  
Ms. Margie Harris, Blood Pattern Section Supervisor  
Ms. Linda Jackson, Controlled Substances Section Chief  
Ms. Katie Jones, Administrative Specialist Forensic Biology  
Mr. Ronald Layne, Director of Administration and Finance  
Ms. Alka Lohmann, Breath Alcohol Section Chief  
Mr. Mike Moore, Questioned Documents Section Chief  
Ms. Carisa Onorato, Administrative Specialist Breath Alcohol  
Mr. Kevin Patrick, Western Laboratory Director  
Ms. Julia Pearson, Toxicology Section Chief  
Mr. James Pickelman, Firearm/Toolmark Section Chief  
Mr. Steven Sigel, Deputy Director  
Mr. Sherwood Stroble, Policy, Planning and Budget Manager  
Ms. Amy Wong, Northern Laboratory Director

Ms. Susan Uremovich, Eastern Laboratory Director  
Mr. Robin Young, Latent Section Chief

Call to Order:

Mr. Fisher called the meeting to order at 9:02 a.m.

Mr. Fisher acknowledged Wanda Adkins as the temporary secretary for the meeting.

Adoption of Agenda:

Mr. Fisher asked if there were any additions or changes to the draft agenda. There were none. There was a motion to adopt the agenda, seconded and adopted without amendment.

Adoption of Minutes:

Mr. Fisher asked if there were any changes that needed to be made to the draft minutes from the August 7, 2007 meeting. There were none. Mr. Smith made a motion to accept the draft minutes, seconded and accepted by unanimous vote.

Director's Report:

Mr. Marone directed members to the information that had been provided on 30-60-90 day workload summary reports by section as of January 1, 2008.

The DNA Section Chief posting has just closed and interviews will be conducted shortly – again  
Minimum Qualifications include:

- Master's Degree or Waiver by ASCLD
- 3 years of Laboratory Experience as a Forensic Nuclear DNA examiner
- QA/QC Experience
- Expert Testimony
- Research and Methods Development

Building update:

- Northern Laboratory – Construction is continuing with an expected move-in date in April of 2009
- Central Laboratory – Administration space in Biotech 8 is projected to be ready in February
- Eastern Laboratory – We have acquired 5,700 square feet with another 15,000 space to be available in late summer on the 5<sup>th</sup> floor for expansion
- Western Laboratory – In the future we hope to be able to acquire additional land.

Dr. Bieber inquired if some of the future Committee meetings could be held in the regional laboratories. Mr. Marone explained that the Code of Virginia requires all meetings to be held in the Richmond area; however he would check to see if there was a possibility of including trips to the regional laboratories for future meetings.

93  
94 Mr. Marone reported on the following grants: 1) NIJ – Forensic Science Training Development  
95 and Delivery Program – development of new training, enhancement of existing training, and  
96 delivery of new and existing forensic science training – no \$\$ amount specified – application due  
97 Feb. 4; 2) NIJ – Solving Cold Cases with DNA – reviewing, investigating violent crime cold  
98 cases that have potential to be solved using DNA – awards not to exceed \$500,000; 3) NIJ -  
99 Social Science Research in Forensic Science – improve the practice of processing of impression  
100 evidence, including fingerprint, tool marks, bite marks, and shoe prints – no dollar amount  
101 specified for individual awards; 4) NIJ - “Research and Development in the Area of Controlled  
102 Substances Detection and Analysis” – We have submitted a concept paper requesting \$50,100.  
103 The title of the proposed project is “Development of a Thin Layer chromatography Method for  
104 the Separation of Enantiomers Using Chiral Mobile Phase Additives.” The project seeks to find  
105 low cost alternatives for separating enantiomers which are controlled differently, such as  
106 dextromethorphan (NCS) and levomethorphan (Schedule II).

107  
108 The DNA/Serology case file review of all 534,000 files have been reviewed and pre-screened.  
109 You have been provided with a flow chart which outlines in very rough fashion the screening  
110 process. The protocol has been completed in draft form and the Governor’s Office is working  
111 with members of the Virginia Bar to approve a working protocol for these cases. A total of 182  
112 cases have been sent to the contractor with 81 waiting to be sent. DFS is continuing to send  
113 cases for testing as they are determined to be eligible and will be attempting to handle persons  
114 currently incarcerated in an expedited fashion to the extent possible.

115  
116 Mr. Bono inquired about the interpretation of the data from these samples? Mr. Marone  
117 explained that the contractor runs the samples and that DFS employees are evaluating all the  
118 data. Dr. Bieber inquired if the Department was tracking the amount of hours consumed on this  
119 project? Dr. Barron responded that it has taken at least 2 years for the files to be reviewed. The  
120 screening of data by the DFS examiners is difficult.

121  
122 DFS has been validating and training on Y-STR technology for several years and expects to put  
123 this type of testing on-line sometime before July of 2008.

124  
125 The Mitochondrial lab staff has received Mito and CODIS training and has ordered servers for  
126 both programs (Mito and CODIS), they should be operational sometime in February. The  
127 manuals are currently being drafted and the laboratory should be online and processing casework  
128 this spring.

129  
130 Blood vial kits – A new kit has been created which should make the process easier to law  
131 enforcement – the kit will include pictograms and instructions that are more precise. In addition,  
132 the Certificate of Blood Withdrawal will have more user-friendly cuts and instructions to clearly  
133 mark the portions which need to be affixed to the blood vial.

134  
135 Breath alcohol instrumentation – The six month evaluation period will end in late February and  
136 we expect to award contract by early March. The first shipment of instruments will be 75 days  
137 after the contract is awarded with the remaining of the instruments coming 150 days after the  
138 contract is awarded.

139  
140 Legislation – the General Assembly convenes on January 9, 2008. DFS will only have some  
141 housekeeping issues on ammunition and sex offender issues in the DNA data bank.  
142

143 Dr. Barron, DFS, gave a presentation addressing a previous request from the Scientific Advisory  
144 Committee to research and review discipline specific certification requirements from relevant  
145 certification bodies; research and review the training guidelines recommended by Scientific  
146 Working Groups (SWG's) and review the individual DFS examiner training programs, criteria  
147 and recommendations of the certification bodies.  
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149 Norah Rudin, Ph.D. and Arthur Eisenberg, Ph.D., DNA Subcommittee, reported to the  
150 committee information requested on contextual bias and “inconclusive” findings. This report will  
151 be attached as an Addendum to these minutes.  
152

153 “Inconclusives” in the first 31 Mary Jane Burton cases - Dr. Rudin reported that the  
154 subcommittee had full access to all the information they needed. Dr. Eisenberg commented that  
155 DFS has done everything in its power to research samples; he felt that DFS had adequately  
156 fulfilled its obligations to the 31 cases. He further stated that DFS has done an admiral job and  
157 Dr. Rudin agreed.  
158

159 Contextual bias – The subcommittee made recommendations for DFS to consider  
160 minimizing the perception of and potential for contextual bias. The Scientific Committee  
161 decided to give DFS time to review and study the recommendations in depth and report back to  
162 the Committee at its next meeting.  
163

164 New Business – Mr. Denio gave a presentation to the Committee and showed a DVD from 60  
165 Minutes on Lead Based Bullet Analysis conducted by the FBI.  
166

167 The next meeting of the Scientific Committee will be held on August 5, 2008 at 9:00 a.m.  
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169 Public Comments – None  
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171 The meeting adjourned at 1:10 p.m.  
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185 ADDENDUM:

186  
187 January 8, 2007

188 Report to the Commonwealth of Virginia Scientific Advisory Committee by the  
189 DNA Subcommittee on confirmation bias and “inconclusive” conclusions  
190 (Norah Rudin, Ph.D. and Artie Eisenberg, Ph.D.)  
191

192 **Subcommittee statement on contextual bias:**

193 Among the many reasons that Forensic DNA analysis has become the gold standard for forensic  
194 science is the relatively discrete nature of the data. For strong, single source samples, a profile  
195 can readily be determined, and is subject to little or no analyst judgment. However, ambiguity  
196 may arise when interpreting more complex samples, such as those containing multiple  
197 contributors, of poor quality (e.g. degraded or inhibited DNA), of low quantity (e.g. contact  
198 samples), or various combinations of these challenging situations. These kinds of samples are  
199 encountered with increasing frequency, as the sensitivity of the technology has increased, and as  
200 law enforcement has become more sophisticated about the kinds of samples they submit for  
201 analysis. Difficult samples are also frequently encountered when reanalyzing historical cases, in  
202 which samples were not collected and preserved using the precautions necessary for DNA  
203 analysis.

204 It is for these types of challenging samples, where the evidence profile may not exactly “match”  
205 a reference profile, that confirmation bias becomes a concern. The interpretation of an  
206 evidentiary DNA profile should not be influenced by information about a subject’s DNA profile.  
207 Each item of evidence must be interpreted independently of other items of evidence or reference  
208 samples. Yet forensic analysts are commonly aware of submitted reference profiles when  
209 interpreting DNA test results, creating the opportunity for confirmatory bias, despite the best  
210 intentions of the analyst. Furthermore, analysts are sometimes exposed to case information, such  
211 as eyewitness identifications or suspect confessions, that may compound an unintentional  
212 confirmatory bias potentially leading to a false inclusion.

213 While it is clear that forensic scientists must be provided with case information in order assist  
214 law enforcement to determine the best and most relevant samples, and to make informed  
215 decisions about the processing of those samples, it is also clear that the analysts who interpret the  
216 data must not inadvertently allow case information, in particular knowledge of reference profiles,  
217 to influence their interpretation of an evidentiary profile. The following general scheme should  
218 serve to minimize the perception of and potential for confirmatory bias:

- 219 1. Data from each evidentiary sample must be interpreted independently by two qualified  
220 analysts
  - 221 a. Both analysts must determine the evidence profile prior to any comparison with  
222 reference profiles.
  - 223 b. Allele calls should be based on defined objective criteria.
  - 224 c. Each analyst must document the specific allele calls, as well as any other notable  
225 characteristics of the profile, such as poor quality, low quantity, or possible  
226 multiple contributors.
  - 227 d. If, upon comparison, discrepancies are noted, the reasons for and resolution of  
228 any such discrepancies must be fully documented.

- 229        2. Reference profiles will be compared to evidentiary profiles only after the evidence  
230        profiles are interpreted and agreement is reached by the two qualified analysts.
- 231            a. The comparison must be performed independently by each analyst.
- 232            b. If, upon comparison, discrepancies are noted, the reasons for and resolution of  
233            any such discrepancies must be fully documented.
- 234        Members of the subcommittee are willing to work with the laboratory to assist in incorporating these safeguards into  
235        the DNA analysis protocol.

236 **Subcommittee statement on “inconclusives” in the first 31 Mary Jane Burton cases**

237 At the August 8, 2007 Forensic Science Board meeting, A motion was passed to request the  
238 Scientific Advisory Committee to “study, report, and make recommendations on the criteria  
239 being used by the lab to report a case as inconclusive in the Mary Jane case file review.”

240 Because the report containing these conclusions is considered part of the Governor’s working  
241 papers, and no separate reports were prepared by the lab, it became a challenge for the  
242 subcommittee to gain access to the information that it was requested to review. Ultimately the  
243 members of the subcommittee were granted access to the document during a visit to the  
244 laboratory that occurred on January 7, 2008.

245 During this visit it was learned that 9 of the original 31 cases had been reported as  
246 “inconclusive.” Of the remaining 22 cases, in 6 instances, the suspect was excluded as a  
247 contributor of the evidence, however in only 2 of those cases did this information provide the  
248 factual basis for an exoneration. In the other 4 exclusions, the evidence either was not relevant or  
249 did not change the facts of the case, and the convictions stood. In 16 cases, the original suspect  
250 was confirmed as a possible contributor of relevant evidence and the convictions stood.

251 As directed, the subcommittee focused on the 9 “inconclusive” cases. Upon reviewing the  
252 Governor’s report, it was found that 4 of these cases were reported as inconclusive because  
253 appropriate reference samples were not available. In these cases, results were obtained for the  
254 evidence samples that could be compared when and if reference samples were obtained. In 5 of  
255 the 9 “inconclusive” cases it was reported that no results were obtained for any of the evidence  
256 samples. The subcommittee requested access to the sample data to independently assess this  
257 conclusion for each of the 5 cases. Mr. Ban and his support staff kindly provided us with full  
258 access to both the case file and electronic data. For 4 of these 5 cases, the subcommittee agreed  
259 that no results were obtained; the gel lanes were effectively blank. However, in one case, some  
260 weak data was visually observed, and corroboration of this data was found in the case file. Mr.  
261 Ban agreed with the subcommittee that, although the data were very weak, and no reliable  
262 comparisons could be performed with any reference samples, that the existence of the data  
263 should have been reported. Specifically, we agreed on the following statement to describe the  
264 results:

265 *The results indicate the presence of a limited amount of male DNA. It is not possible to*  
266 *determine the number of contributors or if female DNA may also be present. Insufficient*  
267 *information exists to perform a meaningful comparison with any reference sample.*

268 The subcommittee felt that it was important to provide this information so that interested parties  
269 could be fully informed that limited genetic material existed that might be tested in more  
270 sensitive systems, such as Y-STRs or mini-STRs.

271 In light of the confusion resulting from the categorization of these 9 cases as “inconclusive”, the  
272 subcommittee suggested a categorization scheme intended to simplify the reporting process and  
273 also to clarify and limit the responsibilities of the laboratory. Once cases are received back from  
274 the contract laboratory, they can be readily categorized as follows:

